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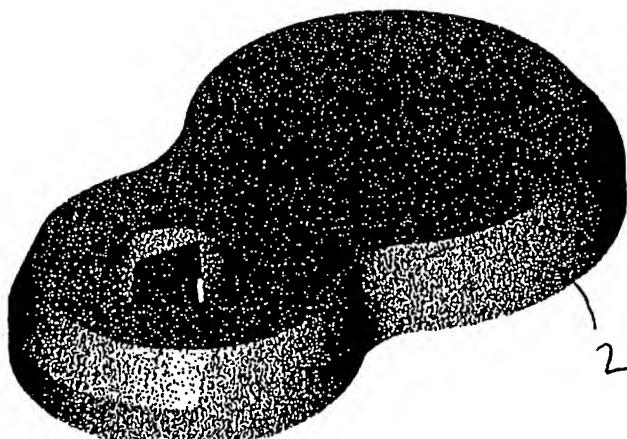
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(54) Title: GASTRIC ACCESS PORT

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adapter or valve is forced open by a cough or gastric pressure build up.

(57) Abstract: An improved mechanism for attaching a feeding tube to a feeding delivery system is presented. The feeding delivery system is attached to an elongated connecting member that is pivotally attached to a bolster. After attachment, the elongated connecting member is rotated to engage another tube at the base of the pivot. Before pivoting, the passageway is blocked by the interference of the elongated connecting member. As the elongated connecting member is rotated, the conduit system aligns and becomes a passageway to and from the stomach. The device provides a three part rigid plastic fit that does not wear out over a number of uses. The device prevents splash events, which occur when a feeding adapter is opened or when the

GASTRIC ACCESS PORT

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FIELD OF THE INVENTION

This invention relates to devices and methods for feeding through gastrostomy feeding or access ports.

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BACKGROUND OF THE INVENTION

- Gastrostomy feeding devices connect the stomach to the exterior of the body through a stoma site. These devices are inserted and left in for the period of time that access to the stomach is needed, and are used as a conduit for feeding, decompression, delivery of medications, and suctioning of fluids. Gastrostomy feeding ports may rise just above the skin level, or they may be an extended tube-like structure with adapters on the exterior end. In either case, an external bolster is used to hold the tube in place to prevent slippage of the external portion of the tube into the stomach.
- Bolsters in the prior art not only provide a mechanism to hold the tube to prevent tube slippage, but also in the tubes that rise just slightly above the skin, these bolsters are charged with connecting to the feeding delivery sets, or syringes used to access the gastrostomy. In this case, a part of the bolster must connect the lumen that traverses the stoma into the stomach with the feeding set or syringe.
- When this connection is opened, reflux of gastric contents is likely.

Prevention of the leakage of gastric contents and the control of gastric pressures in an acidic environment has become a significant design challenge. Materials must be resilient, soft and resealable over time. Unfortunately, many of the gastrostomy feeding port valves or mechanisms for reflux prevention have failed over time, especially after repeated uses.

Valves have been used as a mechanism to block the unwanted flow of gastric contents outward. The prior art includes check valves, such as those available commercially by Novartis under Russo, United States Patent No. 4,944,732. This

product offers a replacement valve which may be screwed into place. The Button, another commercially available device, offers an external rubber flip cap and a tongue valve placed in the stomach end of the communicating lumen. Over time, both the rubber flip cap and the tongue valve become embrittled, and will fail to function.

5 *Copenhaver, et al.* United States Patent No. 5,720,734, commercially available as the Passport from Wilson-Cook Medical, Inc., discusses an "s" slit valve, which lies under compression in a rigid plastic lumen. Because it is a rubber valve, it is fraught with the same embrittlement issues, especially after being violated multiple 10 times per day over a long period.

Prior art devices are formed of rigid plastic and fit only feeding delivery set tips and use an adapter to fit the bolus syringe tips, or have been soft rubber or plastic and subject to breakage upon repeated usage. There is a need for a long term 15 indwelling gastrostomy feeding port which does not utilize a soft rubber valve system or soft rubber flip caps, but which serves to control flow to and from the stomach. This required device eliminates the need for rubber valves and rubber flip caps.

SUMMARY OF THE INVENTION

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A connecting member is housed within a bolster of rigid or soft plastic or rubber. The connecting member pivots about a point which is located at one end of the connecting member. A lumen is connected to the stomach downstream of the pivot point. The lumen is closed when the connecting member is closed and inactive. 25 In the closed or inactive position, the connecting member is substantially perpendicular to the lumen connecting the stomach. The connecting member in the inactive state blocks the flow of the lumen by physically obstructing the flow path.

Before activating the connecting member, the feeding delivery set or syringe is placed into the lumen of the connecting member. This lumen engages the tip of 30 the feeding delivery set tip or syringe as it is the matching taper and provides a plastic-to-plastic fit. To activate, the connecting member is pivoted substantially 90 degrees and the lumen is open. Once the desired procedure of delivering or withdrawing solutions is accomplished, the connecting member is pivoted back into horizontal position and renders the lumen inactive.

By one aspect of the present invention, there is provided a long-term indwelling catheter with an improved anti-reflux mechanism, which remains sealed over multiple uses. The improved anti-reflux mechanism is useful to any feeding gastrostomy system that is left indwelling in a patient and used repeatedly on a daily basis.

The pivoting connecting member may be formed of rigid plastic materials housed in a matrix of a rigid or a resilient bolster connecting the lumen of the feeding tube. The pivoting connecting member is inactive and blocking the lumen of the feeding tube when it is at rest. Upon connection to a syringe or feeding delivery set, 10 the pivoting connecting member is rotated 90 degrees until it is aligned with the feeding lumen. Additional embodiments include the plastic to plastic fit and taper of the pivoting connecting lumen to the feeding delivery sets and to the feeding bolus syringe tip, thus avoiding the use of adapters for engagement.

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DESCRIPTION OF THE DRAWINGS

Figure 1 is a perspective view of the device showing the bottom of the bolster in which the pivoting connecting tube is housed.

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Figure 2 is a perspective view of the device showing the top of bolster in which the pivoting connecting tube is housed.

Figure 3 is an exploded view showing the device with a novel connecting apparatus.

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DETAILED DESCRIPTION OF THE INVENTION

Referring now to the drawing Figures, **Figures 1 and 2** show an anchoring device such as bolster 2 which forms a housing for a connecting member 4. The 30 bolster may be formed of various materials. It is preferred that the bolster be formed of rigid or soft plastic material, or of rubber. The bolster should be formed of materials which are easy to clean, do not readily oxidize, and do not promote the growth of bacteria.

As shown in **Figure 2**, the connecting member is housed within the bolster. One end 8 of the connecting member is preferred to extend beyond the bolster, so that the bolster may be connected to a feeding tube, a syringe or other delivery device, without the necessity of pivoting or rotating the connecting member, so that
5 the device remains in the closed position during connection.

The opposite end of the connecting member is attached to the bolster by means of a pivotal connection. Accordingly, the connecting member may be connected or rotated relative to the bolster. When the connecting member is in the position shown in **Figure 2**, the device is in the closed, or inactive position. That is,
10 access to the stomach through the lumen is blocked by the connecting member.

The lumen is connected to the bottom of the bolster. **Figure 2.** When the connecting member is rotated 90°, so that it is in the position as shown in **Figure 3**, a direct access or opening is available from an upper end of the connecting end, through the connecting member, and into the bolster, through the lower opening of
15 the bolster, and into the lumen. When the connecting member is rotated to the position shown in **Figure 2**, the connecting member blocks the opening to the bolster, also blocking access to the lumen.

As shown, a channel is formed in the bolster. The connecting member is held within the channel, so that the connecting member can be fully rotated 90°. The end
20 of the connecting member extends beyond the end of the bolster. Since the sides of the connecting member are solid and non-porous, as the connecting member is rotated to the position as shown in **Figure 2**, it blocks the opening to the bolster, preventing access to the lumen.

In use, the device is connected to a lumen. A novel method and device for
25 connecting the lumen is discussed hereinafter.

The device is connected to the lumen at the bottom opening of the bolster; an upper end of the connecting member is connected to a feeding tube, syringe, or similar supply conduit. The connecting member is formed to accept appropriate connections. After the feeding tube, syringe or other conduit is connected, the
30 connecting member is rotated to the position shown in **Figure 3**. The device is now in the open or active position, with an unobstructed conduit available from the supply source through the device and into the lumen. At the end of the feeding process or other procedure, the connecting member is rotated from the position shown in

Figure 3, to the position shown in **Figure 2**, and the supply conduit is removed. The device remains in the position as shown in **Figure 2** until access to the stomach is again required for feeding, administering medicines, evacuating the stomach, or the performance of other procedures.

5 The bolster is designed to lie flat against the stomach, so that it is not subject to hooking or snagging or other interaction which could present a hazard to the patient. Similarly, the connecting member is retained within a channel 10 in the bolster, so that the connecting member is not subject to hooking or snagging.

10 A novel means of connecting the lumen is also presented. Most tubes or conduits used with feeding tubes in the prior art are formed of silicone. Silicone is an appropriate material from which form the lumen which is used as the feeding tube. However, certain problems are also present when silicone tubes are used. Silicone tubes tend to have a low coefficient of friction, and are particularly slippery when operating in the wet conditions normally encountered with feeding tubes. Silicone 15 tubes will tend to slip back into the stomach, which can present significant problems for the patient. The tube may not completely slip into the stomach, but it may slip far enough so that the stomach acids are introduced into the body, which can cause sepsis and other problems.

15 In the present invention, a collar 12 is formed having an inside diameter which is approximately the same size as, or perhaps or slightly smaller than, the outside diameter of the resilient tube 14 which is used as the feeding tube. The fit should be such that the collar can be forced onto the tube by manual pressure, but it will not readily slide. The diameter is also such that the collar does not compress to interfere with the flow of material through the tube.

20 The collar should either have threads, or a series of peaks and valleys, or similar irregular shape formed on the inside diameter. For example, a nut of the appropriate inside diameter could be used. Threaded, stainless steel nuts are appropriate candidates for forming the collar. As shown in **Figure 3**, the fitting 16 on the bottom of the device has male threads. The outside diameter of this fitting is 25 preferred to be slightly larger than the inside diameter of the lumen, so that the male threads are engaged with the lumen, and the fitting is advanced by threading it into the lumen.

30 The male threads are used as a mandrel to force the tubing into the threads or other peaks and valleys which are formed in the collar. Threads of the fitting of

the device do not engage the threads of the collar. It is not necessary that the collar be threaded *per se*, it is only necessary that the collar have an irregular surface such as peaks and valleys which will engage and grip the silicone, or other material from which the lumen is formed, when forced against the collar by the male threads.

- 5 Accordingly, the tubing which forms the lumen does not hold the fitting in place by the engagement of the threads, but rather, it is held in place by the forcing of the lumen material against the inside of the collar.

If a longer tube is desired, such as in nursing home use, the longer tube, may be attached to the connecting member of the bolster. If the tube clogs, it may be 10 removed and replaced. In the prior art, clogging is relieved by replacing the entire tube in the endoscopy suite.

The connecting member may be removable. Removal of the connecting member facilitates insertion of laparoscopic or endoscopic instruments. Gastric or esophageal therapy may be performed, and the pivotal connector subsequently 15 replaced. The feeding tube acts as a trocar for laproscopic surgery requiring gastric access.

WHAT IS CLAIMED IS:

1. A device for insertion into a body cavity to selectively transport liquids to and from said body cavity, comprising:
 - a tube comprising a lumen which traverses from an interior of said body cavity to an exterior of said body cavity;
 - an anchoring device that is attached to said tube, wherein said anchoring device comprises a housing, and an elongated connector member having a lumen extending from a first end of said elongated connector member to a second end of said elongated connector member, said elongated connector member being pivotally mounted to said housing, wherein said elongated connecting member is pivoted to align said lumen of said elongated connecting member with said lumen of said tube to facilitate transportation of liquids to and from said body cavity, and said elongated connecting member is alternately pivoted to prevent alignment of said lumen of said elongated connecting member with said lumen of said tube holding mechanism to prevent transportation of liquids to and from said body cavity.
2. A device for insertion into a body cavity to selectively transport liquids to and from said body cavity as described in Claim 1, wherein, when said elongated connecting member is alternately pivoted to prevent alignment of said lumen of said elongated connecting member with said lumen of said tube holding mechanism, said elongated connecting member covers an opening of said lumen of said tube.
3. A device for insertion into a body cavity to selectively transport liquids to and from said body cavity as described in Claim 1, wherein, when said elongated connecting member is alternately pivoted to prevent alignment of said lumen of said elongated connecting member with said lumen of said tube holding mechanism, said elongated connecting member extends beyond an end of said housing.

4. A device for insertion into a body cavity to selectively transport liquids to and from said body cavity as described in Claim 2, wherein, when said elongated connecting member is alternately pivoted to prevent alignment of said lumen of said elongated connecting member with said lumen of said tube holding mechanism and said elongated connecting member covers an opening of said lumen of said tube, said elongated connecting member extends beyond an end of said housing.
5. A device for insertion into a body cavity to selectively transport liquids to and from said body cavity as described in Claim 1 or 2, wherein said housing of said anchoring device comprises a channel therein, and wherein, when said elongated connecting member is alternately pivoted to prevent alignment of said lumen of said elongated connecting member with said lumen of said tube, said elongated connecting member rests in said channel.
6. A device for insertion into a body cavity to selectively transport liquids to and from said body cavity as described in Claim 3 or 4, wherein said housing of said anchoring device comprises a channel therein, and wherein, when said elongated connecting member is alternately pivoted to prevent alignment of said lumen of said elongated connecting member with said lumen of said tube holding mechanism, said elongated connecting member rests in said channel and extends beyond an end of said channel.
7. A device for insertion into a body cavity to selectively transport liquids to and from said body cavity as described in Claim 1, wherein said tube is formed of a resilient material, and further comprising a fitting that extends from a side of said anchoring device that is opposite said elongated connecting member, said fitting having threads that engage an interior of said tube, said tube having a collar positioned on an upper end thereof, said collar having an irregular shape which engages an exterior of said tube, wherein said fitting forces said tube against said collar to hold said tube in place relative to said anchoring device.

8. A device for insertion into a body cavity to selectively transport liquids to and from said body cavity as described in Claim 7, wherein said collar is a nut, and wherein said irregular shape is provided by threads formed in said nut.
9. A device for insertion into a body cavity to selectively transport liquids to and from said body cavity, comprising a tube formed of a resilient material, and further comprising a fitting having threads that engage an interior of said tube, said tube having a collar positioned on a upper end thereof, said collar having an irregular shape which engages an exterior of said tube, wherein said fitting forces said tube against said collar to hold said tube in place.
10. A device for insertion into a body cavity to selectively transport liquids to and from said body cavity as described in Claim 9, wherein said collar is a nut, and wherein said irregular shape is provided by threads formed in said nut.
11. A device for insertion into a body cavity to selectively transport liquids to and from said body cavity as described in Claim 9 or 10, wherein said fitting extends from an anchoring device.
12. A device for insertion into a body cavity to selectively transport liquids to and from said body cavity, comprising:
 - a tube comprising a lumen which traverses from an interior of said body cavity to an exterior of said body cavity;
 - an anchoring device that is attached to said tube, wherein said anchoring device comprises a housing, and an elongated connector member, said elongated connector member being pivotally mounted to said housing, wherein said elongated connecting member is pivoted to align with said lumen of said tube to facilitate transportation of liquids to and from said body cavity, and said elongated connecting member is alternately pivoted to prevent alignment of said elongated connecting member with said lumen of said tube to prevent transportation of liquids to and from said body cavity.

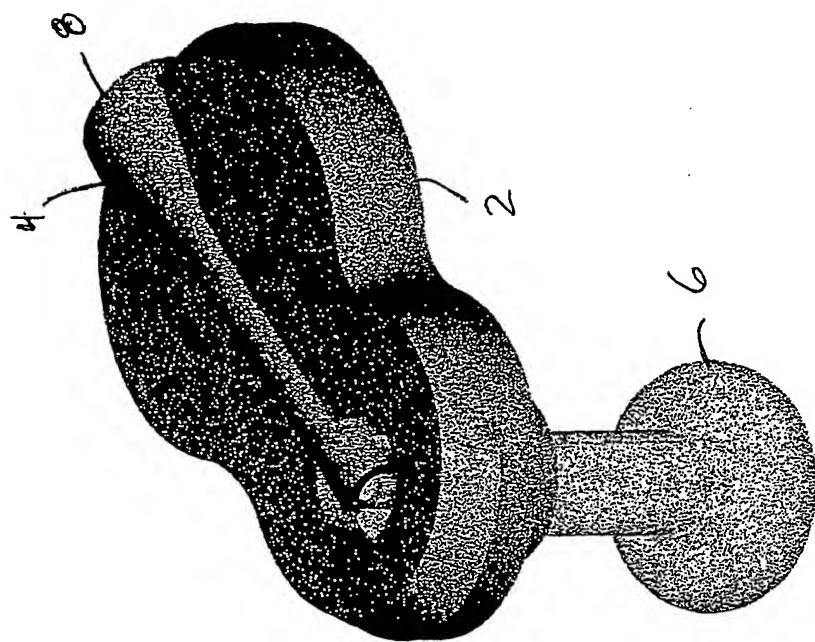


Fig 2

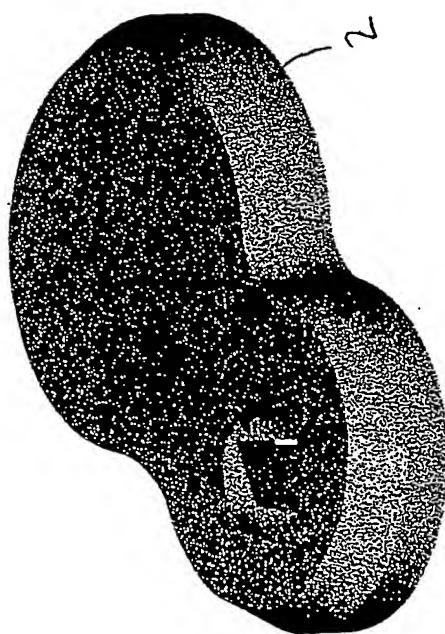


Fig 1

